## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Claims 1-21 (canceled).

22 (currently amended). A method of treating infections an infection of the lower respiratory tract, in a human or animal comprising administering to a patient in need thereof an amount of liposomes sufficient to treat the infection, a pharmaceutical preparation to the lower respiratory tract, said liposomes containing preparation comprising inhalable liposomes combined with povidone iodine.

Claims 23-31 (canceled).

- 32 (currently amended). The method according to claim 22 or 23, wherein the liposomes have a size diameter in the range between from about 1  $\mu$ m and to about 50  $\mu$ m.
- 33 (currently amended). The method according to claim 32, wherein the liposomes have a size diameter in the range between from 20  $\mu$ m to and 30  $\mu$ m diameter for application to the trachea.

Claims 34-35 (canceled).

- 36 (currently amended). The method of claim 22 or 23, wherein the <u>liposomes</u> preparation additionally <u>further contain an</u> comprises at least one anesthetically active agent.
- 37 (currently amended). The method of claim 22 or 23, wherein the <u>liposomes</u> preparation further comprises contain a pharmaceutically acceptable <u>additives</u>.
- 38 (currently amended). The method of claim 22 or 23, wherein the liposomes are suitable for administration is by inhaling the liposomes via nebulization or aerosolization.
- 39 (currently amended). The method of claim 22 or 23, wherein the preparation comprises liposomes are in the form of a tablet, a gelatin capsule, a powder, a spray, an emulsion or a dispersion containing the liposomes and povidone iodine in a pharmaceutically acceptable solid or liquid formulation, which is suitable for the generation of inhalable particles.
- 40 (currently amended). The method of claim 22 32, wherein said preparation comprises:
- (a) comprising a pharmaceutically acceptable liposome membrane forming substance; and

- (b) <u>liposomes contain</u> between 0.1 to 10%, by weight, <u>povidone PVP</u> iodine, wherein the liposomes are in a size between about 1  $\mu$ m and about 50  $\mu$ m.
- 41 (currently amended). The method of claim 40, wherein the liposomes <u>have a diameter</u> in the are in a size range between from 20  $\mu$ m and to 30  $\mu$ m diameter for application to the treachea.

Claims 42-50 (canceled).

51 (currently amended). The method of claim 22 or 23, wherein the liposomes have a diameter size in the range between from about 1  $\mu$ m to and 30  $\mu$ m.

Claims 52-54 (canceled).

- 55 (currently amended). The method of claim 22 or 23, wherein the liposomes have a diameter size in the range between from 10  $\mu$ m and to 20  $\mu$ m diameter for application to the bronchi.
- 56 (currently amended). The method of claim 22, wherein the liposomes have a diameter size in the range between from 1  $\mu$ m and to 6  $\mu$ m diameter for application to the alveoli.
- 57 (currently amended). The method of claim 22, wherein the liposomes have a diameter size in the range between from 2  $\mu$ m and to 5  $\mu$ m diameter for application to the alveoli.
- 58 (currently amended). The <u>method of claim 22</u> preparation of claim 17, wherein said <u>liposomes comprise a</u> liposome membrane forming substance that is present in an amount between 1 to 5%, by weight, of the <u>liposomes preparation</u>.
- 59 (currently amended). The <u>method of claim 22</u> preparation of claim 17, wherein said <u>liposomes are liposome membrane forming substance comprises</u> lecithin <u>liposomes</u>.
- 60 (currently amended). The method of claim 40, wherein said <u>liposomes comprise a</u> liposome membrane forming substance <u>that</u> is present in an <u>amount</u> between about 1 to 5%, by weight, of the <u>preparation liposomes</u>.
- 61 (currently amended). The method of claim 40, wherein said <u>liposomes are liposomes</u> membrane forming substance comprises lecithin <u>liposomes</u>.

Claims 62-63 (canceled).

64 (new). The method of claim 40, wherein said liposomes contain about 0.1% to 2% by weight of povidone iodine.

- 65 (new). The method of claim 22, wherein the infection is in the treachea.
- 66 (new). The method of claim 22, wherein the infection is in the bronchi.
- 67 (new). The method of claim 22, wherein the infection is in the alveoli.
- 68 (new). The method of claim 22, wherein the liposomes further contain a corticosteroid.
- 69 (new). The method of claim 22, wherein the liposomes further contain a second antiseptic agent.
- 70 (new). The method of claim 69, wherein said second antiseptic agent is an oxygen-releasing compound, a halogen-releasing compound, a silver compound, a mercury compound, a formaldehyde-releasing compound, an alcohol, a phenol, a quinoline, an acridine, a hexahydropyrimidine, a quaternary ammonium compound, or a guanidine.
  - 71 (new). The method of claim 22, wherein the liposomes further contain an antibiotic.
- 72 (new). The method of claim 22, wherein said liposomes further contain dexpanthenol, an allantoine, an azulene, a tannine, or a vitamin B compound.
- 73 (new). The method of claim 22, wherein the infection is a bacterial, fungal, or viral infection.
  - 74 (new). The method of claim 22, wherein the patient is human.